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76

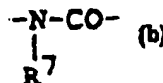
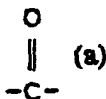
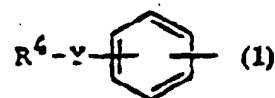
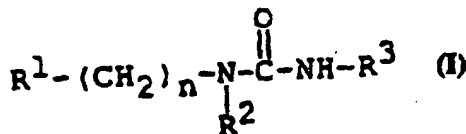
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(71) Applicant (for all designated States except US): FUJISAWA PHARMACEUTICAL CO., LTD. [JP/JP]; 4-7, Doshomachi 3-chome, Chuo-ku, Osaka-shi, Osaka 541 (JP).		Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.	
(72) Inventors; and (75) Inventors/Applicants (for US only): TERASAWA, Takeshi [JP/JP]; 1625-302, Matsugaokanakamachi, Kawachinagano- shi, Osaka 586 (JP). TANAKA, Akira [JP/JP]; 9-10-302, Nakano-cho, Takarazuka-shi, Hyogo 665 (JP). CHIBA, Toshiyuki [JP/JP]; 1-1-503, Nakatsuji-cho, Nara-shi, Nara 630 (JP). TAKASUGI, Hisashi [JP/JP]; 3-116-10, Mozu Umekita, Sakai-shi, Osaka 591 (JP).			

(54) Title: UREA DERIVATIVES AND THEIR USE AS ACAT-INHIBITORS

(57) Abstract

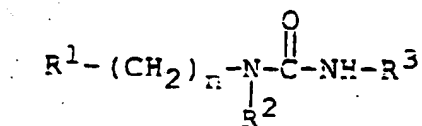
Urea derivatives of formula (I), wherein R¹ is a group of formula (1) (in which R⁴ is aryl which may have suitable substituent(s), or heterocyclic group which may have suitable substituent(s), and Y is bond, lower alkylene, -S-, -O-, (a), -CH-, -CONH-, (b), (in which R² is lower alkyl), -NHSO₂-, -SO₂NH-, -SO₂NHCO- or -CONHSO₂-); or thiazolyl, imidazolyl, pyrazolyl, pyridyl, thienyl, furyl, isoxazolyl or chromanyl, each of which may have suitable substituent(s); R² is lower alkyl, lower alkoxy(lower)alkyl, cycloalkyl, ar(lower)alkyl which may have suitable substituent(s), heterocyclic group or heterocyclic(lower)alkyl, R³ is aryl which may have suitable substituent(s) or heterocyclic group which may have suitable substituent(s), and n is 0 or 1, and a pharmaceutically acceptable salt thereof which are useful as a medicament in the treatment of hypercholesterolemia, hyperlipidemia and atherosclerosis.



- 210 -

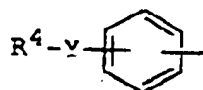
C L A I M S

1. A compound of the formula :



wherein

R^1 is a group of the formula :



(in which

R^4 is aryl which may have suitable substituent(s), or heterocyclic group which may have suitable substituent(s), and

Y is bond, lower alkylene, -S-, -O-, $\overset{\text{C}}{\parallel}$ -C-,
 =CH-, -CONH-, -N-CO-, (in which R^7 is lower alkyl),
 $\underset{\text{R}^7}{\text{N}}$
 -NHSO₂-, -SO₂NH-, -SO₂NHCO- or -CONHSO₂-);
 or

thiazolyl, imidazolyl, pyrazolyl, pyridyl, thienyl, furyl, isoxazolyl or chromanyl, each of which may have suitable substituent(s);

R^2 is lower alkyl, lower alkoxy(lower)alkyl, cycloalkyl, ar(lower)alkyl which may have suitable substituent(s), heterocyclic group or heterocyclic(lower)alkyl,

R^3 is aryl which may have suitable substituent(s) or heterocyclic group which may have suitable

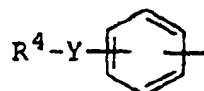
WO 96/10559

PCT/JP95/01982

- 211 -

substituent(s), and
n is 0 or 1,
and a pharmaceutically acceptable salt thereof.

- 5 2. A compound of claim 1, wherein
R¹ is a group of the formula :



(in which

R⁴ is phenyl which may have 1 to 3 substituent(s)
selected from the group consisting of
15 halogen, lower alkyl, di(lower)alkylamino,
protected amino, cyano, heterocyclic group
which may have mono(or di or tri)-
ar(lower)alkyl, hydroxy, protected hydroxy
and mono(or di or tri)halo(lower)alkyl;
20 or thienyl, pyrazolyl, imidazolyl,
triazolyl, pyridyl, pyrrolyl, tetrazolyl,
oxazolyl, thiazolyl, oxadiazolyl,
piperazinyl, thiazolidinyl or
methylenedioxyphenyl, each of which may have
25 1 to 3 substituent(s) selected from the
group consisting of lower alkyl, mono(or di
or tri)ar(lower)alkyl and oxo;

30 Y is bond, lower alkylene, -S-, -O-, -C-, =CH-,
-CONH-, -N-CO- (in which R⁷ is lower alkyl),
-NHSO₂-, -SO₂NH-, -SO₂NHCO- or -CONHSO₂-);
or
35 thiazolyl, imidazolyl, pyrazolyl, pyridyl,